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WHAT IS CLAIMED IS:

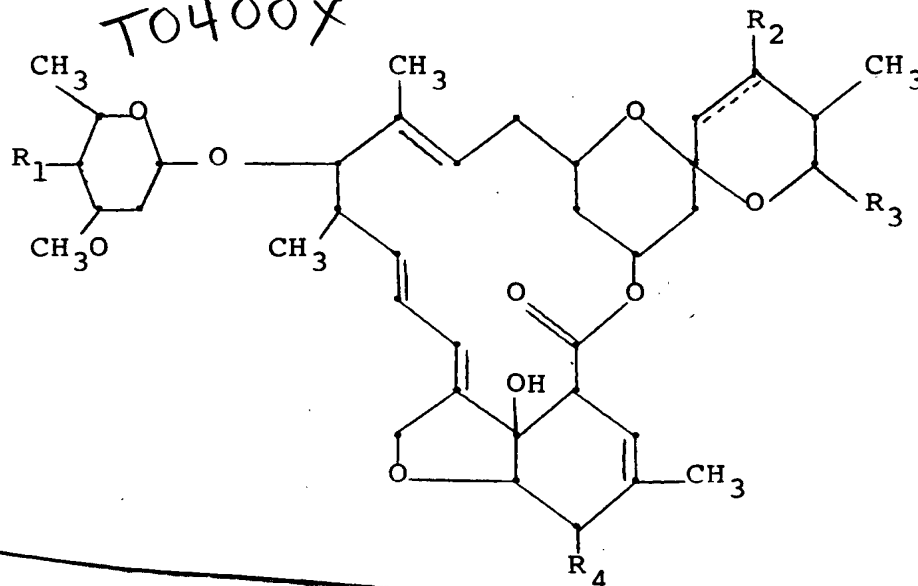
1. A compound having the formula:

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T0400X

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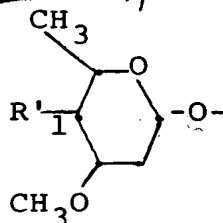
PS

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wherein R_1 is $\equiv O$, $\frac{1}{n}NR_5R_6$ or

B

T0401X



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PS

P₁

wherein $P_1 R'_{140}$ is $\equiv O$ or $\frac{1}{n}NR_5R_6$:

R_5 and R_6 are independently hydrogen, loweralkyl, loweralkanoyl, substituted benzenesulfonyl wherein the substituent is halogen; or loweralkyl sulfonyl;

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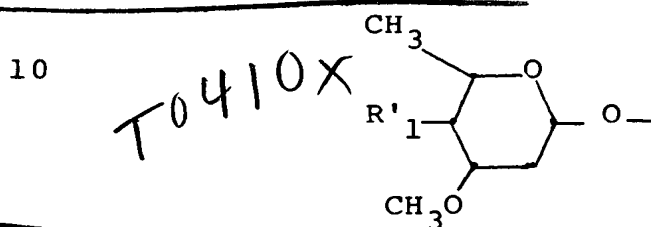
P_1 R_2 is hydrogen or hydroxy;

L R_3 is sec-butyl or iso-propyl;

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P₁
L R₄ is hydroxy or methoxy;
and the broken line indicates a single or a
double bond at the 22,23-position provided that R₂
can only be hydroxy when the broken line indicates a
5 single bond.

2. The compound of Claim 1 wherein R₁ is:



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40, 50 and R'₁ is =O, or NR_5R_6 wherein R₅ and R₆
are independently hydrogen, methyl or acetyl.

50 20 is =O.

3. The compound of Claim 2 wherein R'₁
40 is =O.

4. The compound of Claim 3 which is
41 4"-keto-avermectin Bla/Blb.

25 5. The compound of Claim 3 which is
41 4"-keto-22,23-dihydro avermectin Bla/Blb.

30 6. The compound of Claim 2 wherein R'₁ is
41 NR_5R_6 and R₅ and R₆ are independently
hydrogen, methyl or acetyl.

7. The compound of Claim 6 which is
41 4"-deoxy-4"-amino-avermectin Bla or Blb.

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8. The compound of Claim 6 which is
(41) 4"-deoxy-4"-amino-22,23-dihydro-avermectin Bla or Blb.

9. The compound of Claim 6 which is
(41) 5 4"-deoxy-4"-acetylamino-avermectin Bla or Blb.

10. The compound of Claim 6 which is
(41) 4"-deoxy-4"-acetylamino-22,23-dihydro avermectin Bla
or Blb.

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11. The compound of Claim 6 which is
(41) 4"-deoxy-4"-N,N-dimethylamino-avermectin Bla or Blb.

12. The compound of Claim 6 which is
(41) 15 4"-deoxy-4"-N,N-dimethylamino-22,23-dihydro
avermectin Bla or Blb.

N
K
N
P
20 13. A process for the preparation of the
compounds of Claim 1 wherein R_1 or R'_1 is =O
which comprises treating a compound wherein R_1 or
 R'_1 is hydroxy with dimethyl sulfoxide and oxalyl
chloride or trifluoroacetic anhydride, or with
N-chlorosuccinimide and dimethylsulfide.

25 14. A process for the preparation of
a compound of Claim 1 wherein R_1 or R'_1 is
-NR₅R₆ wherein R₅ and R₆ are as defined in
Claim 1 which comprises treating the corresponding
compound wherein R_1 or R'_1 is =O with an ammonium
30 salt or methyl ammonium salt and a reducing agent and
optionally treating the thus produced 4"-deoxy-4"-
amino-or 4"-deoxy-4'-amino compound with an acylating

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agent in the presence of a base to produce the 4"-deoxy-4"-acylamino and 4'-deoxy-4'-acylamino compound or optionally treating the amino compound with aqueous formaldehyde and a reducing agent under acidic conditions to produce the corresponding 4"-deoxy-4"-N,N-dimethylamino and 4'-deoxy-4'-N,N-dimethyl-amino compounds.

15. The process of Claim 14 wherein the salts used are the acetate, propionate or benzoate, and the reducing agent is sodium cyano borohydride.

16. The process of Claim 15 wherein the salt used is the acetate.

13. (13) A method for the treatment of ^{helminthiasis} parasitic infections which comprises administering to an animal infected with ^{helminths} parasites, an effective amount of a compound of Claim 1.

14. (14) A composition useful for treating animals infected with ^{helminths} parasites, which comprises an ^{inert} inner carrier and ^{an effective amount of} a compound of Claim 1.

19. A method for the treatment of bacterial infections which comprises administering to an animal infected by bacteria, an effective amount of a compound of Claim 1 wherein R_1 or R'_1 is $-NR_5R_6$ and R_5 and R_6 are as defined in Claim 1.

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20. A composition useful for treating
bacterial infections which comprises an inert carrier
and a compound of Claim 1 wherein R_1 or R'_1 is
-NR₅R₆ and R₅ and R₆ are as defined in
Claim 1.

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